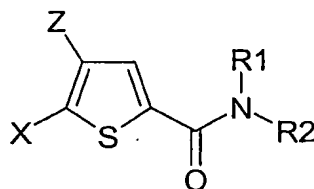


What is claimed is:

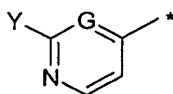
1. A compound of Formula (I) or a salt, solvate, or physiologically functional derivative thereof:



(I)

wherein:

- R1 is hydrogen or C₁₋₆alkyl;
- R2 is selected from the group consisting of C₁₋₆alkyl, C₁₋₄alkylNR⁷R⁸ (wherein R⁷ and R⁸ are independently H or C₁₋₄alkyl), aryl, CH(CH₂OH)aryl, arylC₁₋₆alkyl, aryloxyC₁₋₆alkyl, heteroaryl, heteroarylC₁₋₆alkyl, heterocyclyl and heterocyclylC₁₋₆alkyl, wherein in each case the aryl, heteroaryl or heterocyclyl moiety is optionally substituted by one to five groups selected from the group consisting of halogen, NH₂, hydroxy, cyano, C₁₋₄alkyl, -OCH₂O-, C₁₋₄alkoxy, haloC₁₋₄alkyl, haloC₁₋₄alkoxy, aryl, aryloxy, C₁₋₄alkoxycarbonyl, C₁₋₄hydroxyalkyl, C₁₋₄alkanoyl, C₁₋₄alkylsulfonyl, (CH₂)₀₋₄NHCOOC₁₋₄alkyl, and a group R₃R₄NSO₂ (wherein R₃ and R₄ are independently hydrogen or C₁₋₄alkyl) and a 5- or 6-membered heteroaryl group;
- or R1 and R2, together with the nitrogen atom to which they are joined, form a 5- or 6-membered monocyclic heterocyclic ring or a 9- or 10-membered bicyclic heterocyclic ring wherein at least the ring which contains the nitrogen atom to which R1 and R2 are joined is non-aromatic, and wherein the 5- or 6-membered monocyclic heterocyclic ring or the 9- or 10-membered bicyclic heterocyclic ring is optionally substituted by one to four groups selected from the group consisting of halogen, hydroxy, cyano, C₁₋₄alkanoyl, oxo, C₁₋₄alkyl, C₁₋₄alkoxy, haloC₁₋₄alkyl, haloC₁₋₄alkoxy, aryl, aryloxy and C₁₋₄alkoxycarbonyl;
- X is indazolyl, pyrazolyl or a group



wherein

G is CH or N; and

Y is hydrogen or a group NR₅R₆ (wherein R₅ and R₆ are independently hydrogen, C₁₋₆alkyl), (CH₂)₀₋₆phenyl (wherein the phenyl group is optionally substituted by halogen or OC₁₋₄alkyl);

and

- Z is hydrogen, halogen, cyano or a 5- or 6-membered heteroaryl.

2. A compound as claimed in claim 1, wherein R1 is hydrogen.

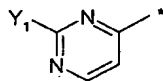
3. A compound as claimed in claim 1 or claim 2, wherein R2 is arylC₁₋₆alkyl optionally substituted by one or two groups selected from the group consisting of halogen, hydroxy, C₁₋₄alkyl, C₁₋₄alkoxy, haloC₁₋₄alkyl, haloC₁₋₄alkoxy, thiadiazolyl and a group R₃R₄NSO₂ wherein R₃ and R₄ are independently hydrogen or C₁₋₄alkyl.

4. A compound as claimed in claim 1 or claim 2, wherein R1 and R2, together with the nitrogen atom to which they are joined, form a 6-membered monocyclic heterocyclic ring or a 10-membered bicyclic heterocyclic ring wherein at least the ring which each contains the nitrogen atom to which R1 and R2 are joined is non-aromatic, wherein the 6-membered monocyclic heterocyclic ring or 10-membered bicyclic heterocyclic ring are both optionally substituted by one or two groups selected from oxo, C₁₋₄alkyl, phenyl and C₁₋₄alkoxycarbonyl.

5. A compound as claimed in any of claims 1-4, wherein X is indazolyl or pyrazolyl.

6. A compound as claimed in any of claims 1-4, wherein X is 4-pyridinyl and Y is hydrogen.

7. A compound as claimed in any of claims 1-4, wherein X is a group:



wherein Y₁ is a group NR₅R₆ wherein R₅ and R₆ are independently hydrogen or C₁₋₆alkyl.

8. A compound as claimed in any of claims 1-8 wherein Z is hydrogen or halogen.

9. A compound as claimed in claim 1, which is:

N-(2-phenylethyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
 N-(3-methoxybenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
 5-(4-pyridinyl)-N-(2-pyridinylmethyl)-2-thiophenecarboxamide
 N-(1-naphthylmethyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
 N-(2-ethoxybenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
 N-(2-bromobenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
 N-(2-fluorobenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
 N-(2-chlorobenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide

N-(2-methylbenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(2-trifluoromethylbenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(2-trifluoromethoxybenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(3-trifluoromethylbenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(3-fluorobenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(3-chlorobenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(3-bromobenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(3-iodobenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(3-methylbenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(3-methoxybenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(3-trifluoromethoxybenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(3-phenoxybenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(4-fluorobenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(4-bromobenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(4-iodobenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(4-trifluoromethylbenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(4-methylbenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(4-trifluoromethoxybenzyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[4-(aminosulfonyl)benzyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[4-(methylsulfonyl)benzyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
5-(4-pyridinyl)-N-(4-pyridinylmethyl)-2-thiophenecarboxamide
N-benzyl-N-methyl-5-(4-pyridinyl)-2-thiophenecarboxamide
5-(4-pyridinyl)-N-[4-(1,2,3-thiadiazol-4-yl)benzyl]-2-thiophenecarboxamide
5-(4-pyridinyl)-N-(3-pyridinylmethyl)-2-thiophenecarboxamide
N-[2-(2-methylphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(3-methylphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(4-methylphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(2-fluorophenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(3-fluorophenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(4-fluorophenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(2-methoxyphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(3-methoxyphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(4-methoxyphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(2-chlorophenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(3-chlorophenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(4-chlorophenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(2-ethoxyphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(3-ethoxyphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(4-ethoxyphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(2-bromophenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(3-bromophenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide

N-[2-(4-bromophenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(2-phenoxyphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(4-phenoxyphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(4-hydroxyphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(3-trifluoromethylphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-[4-(aminosulfonyl)phenyl]ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
2-[[5-(4-pyridinyl)-2-thienyl]carbonyl]-1,2,3,4-tetrahydroisoquinoline
5-(4-pyridinyl)-N-[2-(3-pyridinyl)ethyl]-2-thiophenecarboxamide
5-(4-pyridinyl)-N-[2-(4-pyridinyl)ethyl]-2-thiophenecarboxamide
N-(2-phenoxyethyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(1-piperidinyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[2-(4-morpholinyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
1-phenyl-4-[[5-(4-pyridinyl)-2-thienyl]carbonyl]piperazine
N-(1H-indazol-5-yl)-5-(4-pyridinyl)-2-thiophenecarboxamide
1-phenyl-8-[[5-(4-pyridinyl)-2-thienyl]carbonyl]-1,3,8-triazaspiro[4.5]decan-4-one
ethyl 4-[[[5-(4-pyridinyl)-2-thienyl]carbonyl]amino]-1-piperidinecarboxylate
ethyl 1-[[5-(4-pyridinyl)-2-thienyl]carbonyl]-4-piperidinecarboxylate
N-(1H-benzimidazol-2-ylmethyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
5-(4-pyridinyl)-N-[2-(2-pyridinyl)ethyl]-2-thiophenecarboxamide
N-[2-(3-hydroxyphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[(1R)-1-phenylethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[(1S)-1-phenylethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[(1R)-1-(3-methoxyphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-[(1S)-1-(3-methoxyphenyl)ethyl]-5-(4-pyridinyl)-2-thiophenecarboxamide
N-isopropyl-5-(4-pyridinyl)-2-thiophenecarboxamide
1-methyl-4-[[5-(4-pyridinyl)-2-thienyl]carbonyl]piperazine
N-phenyl-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(2-methoxyphenyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(2-chlorophenyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(4-methoxyphenyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-(4-chlorophenyl)-5-(4-pyridinyl)-2-thiophenecarboxamide
N-benzyl-5-(4-pyridinyl)-2-thiophenecarboxamide
5-(2-amino-4-pyrimidinyl)-N-(3-methoxybenzyl)-2-thiophenecarboxamide
5-(2-amino-4-pyrimidinyl)-N-benzyl-2-thiophenecarboxamide
5-(4-pyrimidinyl)-N-(3-methoxybenzyl)-2-thiophenecarboxamide
5-(1H-indazol-5-yl)-N-(3-methoxybenzyl)-2-thiophenecarboxamide
5-(6-amino-4-pyrimidinyl)-N-(3-methoxybenzyl)-2-thiophenecarboxamide
N-benzyl-4-bromo-5-(4-pyridinyl)-2-thiophenecarboxamide
N-benzyl-4,5-di(4-pyridinyl)-2-thiophenecarboxamide
N-(3-methoxybenzyl)-4,5-di(4-pyridinyl)-2-thiophenecarboxamide
N-benzyl-5-[2-(methylamino)-4-pyrimidinyl]-2-thiophenecarboxamide

N-benzyl-5-(1H-pyrazol-4-yl)-2-thiophenecarboxamide

N-(3-methoxybenzyl)-5-(1H-pyrazol-4-yl)-2-thiophenecarboxamide

or a salt, solvate or physiologically functional derivative thereof.

10. A compound as claimed in any of claims 1-9 for use in therapy.
11. A compound as claimed in any of claims 1-9 for use in the treatment of a disorder mediated by inappropriate ROCK-1 activity.
12. A method of treating a disorder in a mammal, said disorder being mediated by inappropriate ROCK-1 activity, comprising: administering to said mammal a therapeutically effective amount of a compound as defined in any of claims 1-9.
13. Use of a compound as defined in any of claims 1-9 in the preparation of a medicament for use in the treatment of a disorder mediated by inappropriate ROCK-1 activity.
14. A pharmaceutical composition comprising a therapeutically effective amount of a compound as defined in any of claims 1-9 and one or more of pharmaceutically acceptable carriers, diluents and excipients.